## WHAT IS CLAIMED IS:

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1. A composition for fabricating phase-change-material microcapsule, comprising:

5% to 40 % weight percentage concentration of waterborne polyurethane aqueous solution;

phase-change-material;

lipophilic monomer; and

solid wax, wherein the weight percentage concentration of the lipophilic monomer solving in the phase change material is between about 3% and 12%, and the weight ratio of lipophilic monomer to waterborne polyurethane is between about 25% and 50%.

- 2. The composition of claim 1, wherein the waterborne polyurethane in the waterborne polyurethane aqueous solution is selected from a group consisting waterborne polyurethane, 2,2-bis (hydroxymethyl) propionic acid triethylamine salt, diamine containing sulfonate salt and a combination thereof.
- 3. The composition of claim 1, wherein the phase-change-material is an organic compound with polarity.
  - 4. The composition of claim 1, wherein the phase-change-material is a carboxylic ester.

- 5. The composition of claim 4, wherein a carboxylate of the carboxylic ester is selected from a group of formate, acetate and propionate.
- 6. The composition of claim 4, wherein carbon atom number of an alkoxyl of the carboxylic ester are between 10 and 18.
  - 7. The composition of claim 1, wherein the lipophilic monomer is melamine or isocyanate salt.
- 8. The composition of claim 1, wherein the preferred weight ratio of waterborne polyurethane to microcapsule composition is between about 10% and 30%.
- 9. The composition of claim 1, wherein the preferred weight percentage concentration of the lipophilic monomer solving in the phase change material is between about 5% and 10%.
  - 10. The composition of claim 1, wherein the preferred weight ratio of lipophilic monomer to waterborne polyurethane is between about 30% and 45%.
  - 11. A method using the composition of claim 1 for fabricating phase-change-material microcapsule dispersing in a water phase, comprising:
    - putting the composition in a reactor, wherein the composition comprising: the waterborne polyurethane aqueous solution;
- 25 the phase-change-material;

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## the lipophilic monomer; and

the solid wax; emulsify the composition by stirring;

performing at least two stages heating process to elevate a temperature of the emulsified composition; and

adding at least one stabilizer.

12. The method of claim 11, wherein a speed of the emulsify by stirring is between about 4000 rpm and 9000 rpm.

13. The method of claim 11, wherein a time for the emulsion by stirring is

between about 2 minutes and 5 minutes.

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- 14. The method of claim 11, wherein the temperature range is betweenabout 20 degree Celsius and 90 degree Celsius.
  - 15. The method of claim 11, wherein the elevating temperature further comprising:

keeping aconstant temperature at each stage, wherein the duration is from 1 hour to 5 hours at the stage.

16. The method of claim 11, wherein the waterborne polyurethane in the waterborne polyurethane aqueous solution is selected from a group consisting of waterborne polyurethane, 2,2-bis (hydroxymethyl) propionic acid and its triethylamine salt, diamine containing sulfonate salt and a combination thereof.

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- 17. The method of claim 11, wherein the stabilizer is sorbitan monooleate or sodium dodecyl sulfonate.
- 18. The method of claim 11, wherein the phase-change-material is an organic compound with polarity.
  - 19. The method of claim 11, wherein the phase-change-material is a carboxylic ester.

20. The method of claim 19, wherein a carboxylate of the carboxylic ester is selected from a group consisting of formate, acetate and propionate.

- 21. The method of claim 19, wherein carbon atom number of an alkoxyl of the carboxylic ester is between 10 and 18.
  - 22. A phase-change-material for fabricating a microcapsule used between minus 20 degree Celsius and 80 degree Celsius, comprising:

a carboxylic ester, wherein a carboxylate of the carboxylic ester is selected from a group formate, acetate and propionate and carbon atom number of an alkoxyl of the carboxylic ester are between 10 and 28.

23. The phase-change-material of claim 21, wherein carbon atom number of an alkoxyl of the carboxylic ester is preferred between 10 and 18.

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24. The phase-change-material of claim 21, wherein the microcapsule is preferred used between minus 20 degree Celsius and 80 degree Celsius.